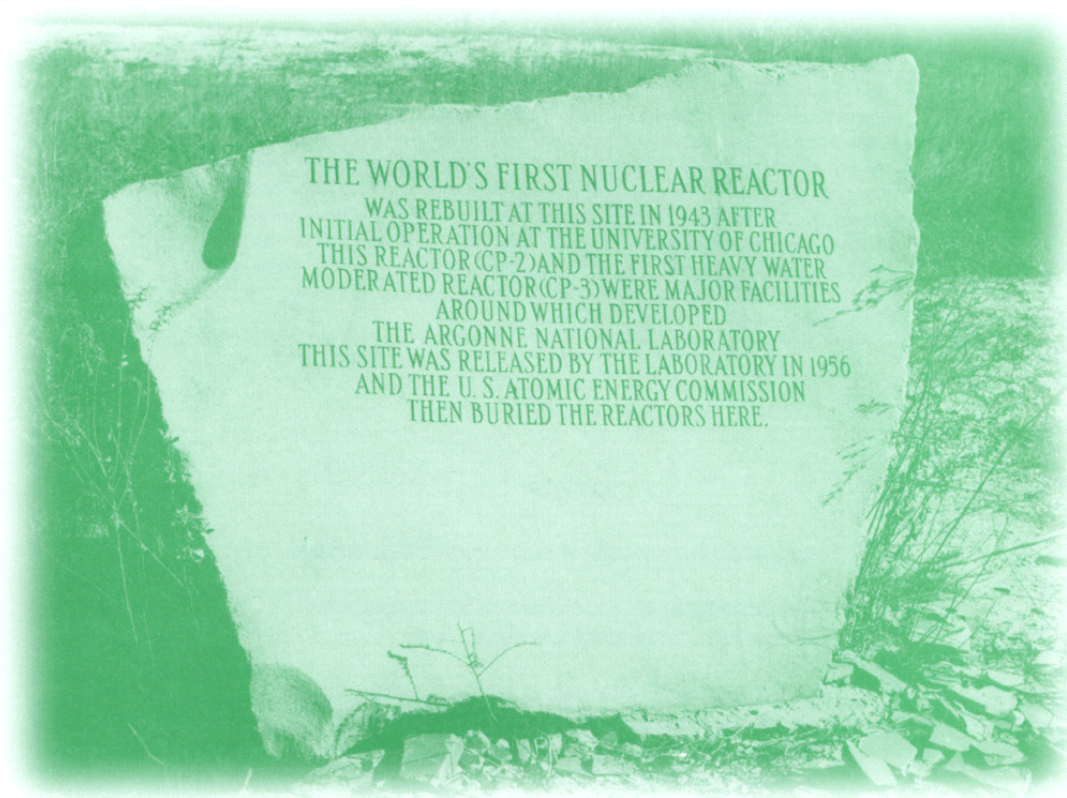


Surveillance of Site A and Plot M

Report for 2000

Norbert W. Golchert
The Office of ESH/QA Oversight



Argonne National Laboratory, Argonne, Illinois 60439
operated by The University of Chicago
for the United States Department of Energy under Contract W-31-109-Eng-38

Argonne National Laboratory, with facilities in the states of Illinois and Idaho, is owned by the United States Government and operated by The University of Chicago under the provisions of a contract with the Department of Energy.

DISCLAIMER

This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency thereof, nor The University of Chicago, nor any of their employees or officers, makes any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof. The views and opinions of document authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof, Argonne National Laboratory, or The University of Chicago.

Available electronically at <http://www.doe.gov/bridge>

Available for a processing fee to U.S. Department of Energy and its contractors, in paper, from:

U.S. Department of Energy
Office of Scientific and Technical Information
P.O. Box 62
Oak Ridge, TN 37831-0062
phone: (865) 576-8401
fax: (865) 576-5728
email: reports@adonis.osti.gov

ANL- 01/1

ARGONNE NATIONAL LABORATORY-EAST
9700 South Cass Avenue
Argonne, Illinois 60439

SURVEILLANCE OF SITE A AND PLOT M

Report for 2000

by

Norbert W. Golchert

The Office of ESH/QA Oversight

May 2001

Prepared for the U. S. Department of Energy
Work Package ANL 035 00 01 -- WBS No. AH-10-05-121I

PREFACE

This report is prepared for the U. S. Department of Energy (DOE) by the The Office of ESH/QA Oversight (EQO) at Argonne National Laboratory-East (ANL-E). The results of the environmental monitoring program at Site A and Plot M and an assessment of the impact of the site on the environment and the public are presented in this publication. Funding to support this program was provided by the Office of Environmental Management through the U. S. Department of Energy Grand Junction Office. This report and some earlier issues of the annual reports are available on the Internet at <http://www.anl.gov/ESH/sitea/2000>.

Most of the tables and some of the figures were prepared by Jennifer Tucker of the Data Management Team. Sample collection and field measurements were conducted under the direction of Ronald Kolzow of the Monitoring and Surveillance Group by:

Michael Cole

Dan Milinko

The analytical separations and measurements were conducted by the Dosimetry and Analytical Services Section of the Chemical Technology Division (CMT) by:

Tim Branch

Denise Seeman

Theresa Davis

Christos Stamoudis

Alan Demkovich

Emmer Thompson

Gary Griffin

John Zhang

Richard Kasper

This manuscript was typed and prepared for publication by Rita M. Beaver (EQO).

TABLE OF CONTENTS

	<u>Page</u>
PREFACE	iii
TABLE OF CONTENTS	v
LIST OF TABLES	vii
LIST OF FIGURES	ix
ABSTRACT	xi
1.0 INTRODUCTION	1-1
1.1 Site History	1-1
1.2 Site Characteristics	1-5
2.0 SUMMARY	2-1
3.0 MONITORING PROGRAM	3-1
3.1 Surface Water	3-2
3.2 Subsurface Water	3-5
3.2.1 Borehole Water - Plot M	3-5
3.2.2 Borehole Water - Site A	3-10
3.2.3 Dolomite Hole Water	3-32
3.2.4 Well Water	3-37
4.0 SUMMARY OF POTENTIAL RADIATION DOSE AND RISK ESTIMATES	4-1
4.1 Dose Estimates	4-1
4.2 Risk Estimates	4-1
5.0 REFERENCES	5-1

TABLE OF CONTENTS (Contd.)

	<u>Page</u>
6.0 APPENDICES	6-1
6.1 Quality Assurance Program	6-1
6.2 Applicable Standards	6-2
6.3 Analytical Methods	6-2
Distribution for ANL-01/1	6-3

LIST OF TABLES

<u>No.</u>	<u>Title</u>	<u>Page</u>
3.1	Tritiated Water Content of Stream Next to Plot M, 2000	3-4
3.2	Tritium in Plot M Borehole Water, 2000	3-7
3.3	Water Level Measurements in Boreholes Near Plot M, 2000	3-8
3.4	Strontium-90 Analyses of Borehole Water Samples Near Plot M, 2000	3-11
3.5	Tritium in Site A Borehole Water, 2000	3-13
3.6	Water Level Measurements in Boreholes Near Site A, 2000	3-14
3.7	Strontium-90 in Site A Borehole Water Samples, 2000	3-15
3.8	Chemical Constituents in Site A Borehole 41, 2000	3-17
3.9	Chemical Constituents in Site A Borehole 42, 2000	3-18
3.10	Chemical Constituents in Site A Borehole 44, 2000	3-19
3.11	Chemical Constituents in Site A Borehole 45, 2000	3-20
3.12	Chemical Constituents in Site A Borehole 46, 2000	3-21
3.13	Chemical Constituents in Site A Borehole 47, 2000	3-22
3.14	Chemical Constituents in Site A Borehole 48, 2000	3-23
3.15	Chemical Constituents in Site A Borehole 49, 2000	3-24
3.16	Chemical Constituents in Site A Borehole 50, 2000	3-25
3.17	Chemical Constituents in Site A Borehole 51, 2000	3-26
3.18	Chemical Constituents in Site A Borehole 52, 2000	3-27
3.19	Chemical Constituents in Site A Borehole 53, 2000	3-28
3.20	Chemical Constituents in Site A Borehole 54, 2000	3-29
3.21	Chemical Constituents in Site A Borehole 55, 2000	3-30
3.22	Chemical Constituents in Site A Borehole 56, 2000	3-31

LIST OF TABLES (Contd.)

<u>No.</u>	<u>Title</u>	<u>Page</u>
3.23	Tritium in Site A Dolomite Holes, 2000	3-34
3.24	Water Level Measurements in Dolomite Holes, 2000	3-35
3.25	Tritiated Water Content of Wells Near Site A/Plot M, 2000	3-38
3.26	Annual Maximum and Average Tritium Concentrations in the Red Gate Woods Wells	3-40
3.27	Uranium Content of Well Water Near Plot M, 2000	3-43
3.28	Inorganic Constituents in Red Gate Woods North Well Water (#5160), 2000	3-44
4.1	Dose From Continuous Exposure to Tritium at Selected Locations, 2000	4-2
4.2	Estimates of Tritium Exposures to a Casual Visitor to Plot M, 2000	4-3
4.3	Annual Average Dose Equivalent in the U. S. Population	4-4
4.4	Risk of Death From Natural Events	4-5

LIST OF FIGURES

<u>No.</u>	<u>Title</u>	<u>Page</u>
1.1	Location of Palos Forest Preserve on Chicago-Area Map	1-2
1.2	Palos Forest Preserve Showing Location of Site A/Plot M Dolomite Holes and Picnic Wells	1-3
3.1	Surface Water Sampling Locations Near Plot M	3-3
3.2	Map of Plot M Palos Site Showing Topography, Intermittent Stream, and Borehole Locations	3-6
3.3	Monitoring Wells at Site A	3-12
3.4	Locations of Dolomite Holes North of Plot M	3-33
3.5	Tritiated Water Concentrations in Red Gate Woods (#5167), Opposite Red Gate Woods (#5159), and Red Gate Woods North (#5160) Wells From 1992 Through 2000	3-41

SURVEILLANCE OF SITE A AND PLOT M
Report for 2000
by
N. W. Golchert

ABSTRACT

The results of the environmental surveillance program conducted at Site A/Plot M in the Palos Forest Preserve area for Calendar Year 2000 are presented. Based on the results of the 1976-1978 radiological characterization of the site, a determination was made that a surveillance program be established. The characterization study determined that very low levels of hydrogen-3 (as tritiated water) had migrated from the burial ground and were present in two nearby hand-pumped picnic wells. The current surveillance program consists of sample collection and analysis of surface and subsurface water. The results of the analyses are used to 1) monitor the migration pathway of water from the burial ground (Plot M) to the handpumped picnic wells, 2) establish if buried radionuclides other than hydrogen-3 have migrated, and 3) generally characterize the radiological environment of the area. Hydrogen-3 in the Red Gate Woods picnic wells was still detected this year, but the average and maximum concentrations were significantly less than found earlier. Tritiated water continues to be detected in a number of wells, boreholes, dolomite holes, and a surface stream. Analyses since 1984 have indicated the presence of low levels of strontium-90 in water from a number of boreholes next to Plot M. The available data does not allow a firm conclusion as to whether the presence of this nuclide represents migration prior to the installation of the boreholes or movement that may have occurred before Plot M was capped. The results of the surveillance program continue to indicate that the radioactivity remaining at Site A/Plot M does not endanger the health or safety of the public visiting the site, using the picnic area, or living in the vicinity.

INTRODUCTION

1.1 Site History

This report presents and discusses the surveillance data obtained during 2000. The surveillance program is the ongoing activity that resulted from the 1976-1978 radiological characterization of the former site of Argonne National Laboratory and its predecessor, the University of Chicago's Metallurgical Laboratory. This site was part of the World War II Manhattan Engineer District Project and was located in the Palos Forest Preserve southwest of Chicago, IL. The Laboratory used two locations in the Palos Forest Preserve: Site A, a 19-acre area that contained experimental laboratories and nuclear reactor facilities; and Plot M, a 150 ft x 140 ft area used for the burial of radioactive waste. These locations are shown in Figure 1.1 and Figure 1.2. Previous comprehensive reports on this subject,^{1,2} provide additional detail and illustrations on sampling locations and provide descriptive material along with the results through 1981. There are annual reports for 1982 through 1999.³⁻²⁰ While earlier data will not be repeated in this report, reference is made to some of the results.

Operations at Site A began in 1943 and ceased in 1954. Among the research programs carried out at Site A were reactor physics studies, fission product separations, tritium recovery from irradiated lithium, and work related to the metabolism of radionuclides in laboratory animals. Radioactive waste and radioactively-contaminated laboratory articles from these studies were buried at Plot M. At the termination of the programs, the reactor fuel and heavy water, used for neutron moderation and reactor cooling, were removed and shipped to Oak Ridge National Laboratory. The biological shield for the CP-3 reactor located at Site A, together with various pipes, valves, and building debris, was buried in place in 1956.

Burial of radioactive waste at Plot M began in 1944 and was discontinued in 1949. Waste was buried in six-foot deep trenches and covered with soil until 1948, after which, burial took place in steel bins. The steel bins were removed in 1949 and sent to Oak Ridge National

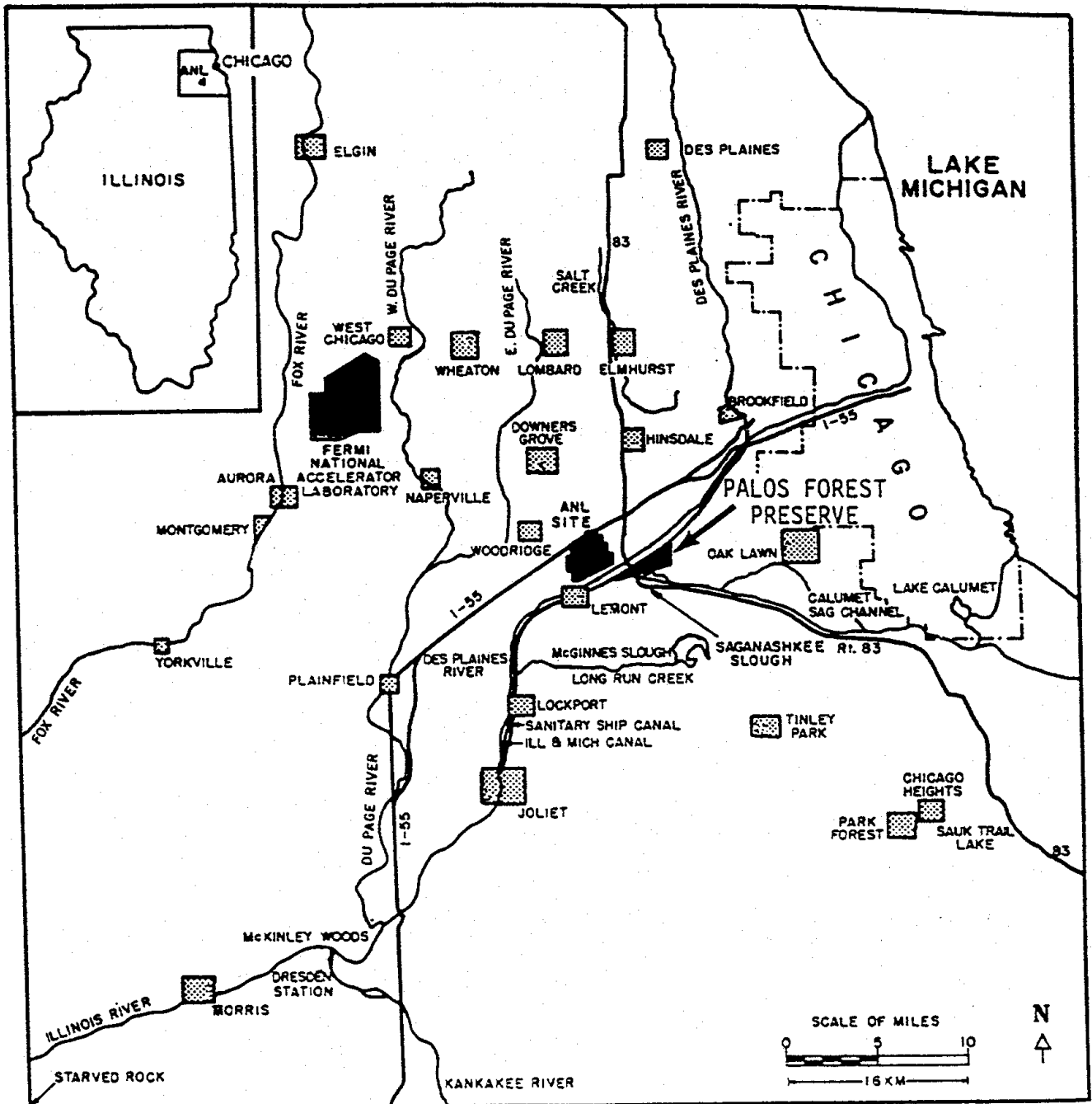


Figure 1.1 Location of Palos Forest Preserve on Chicago-Area Map

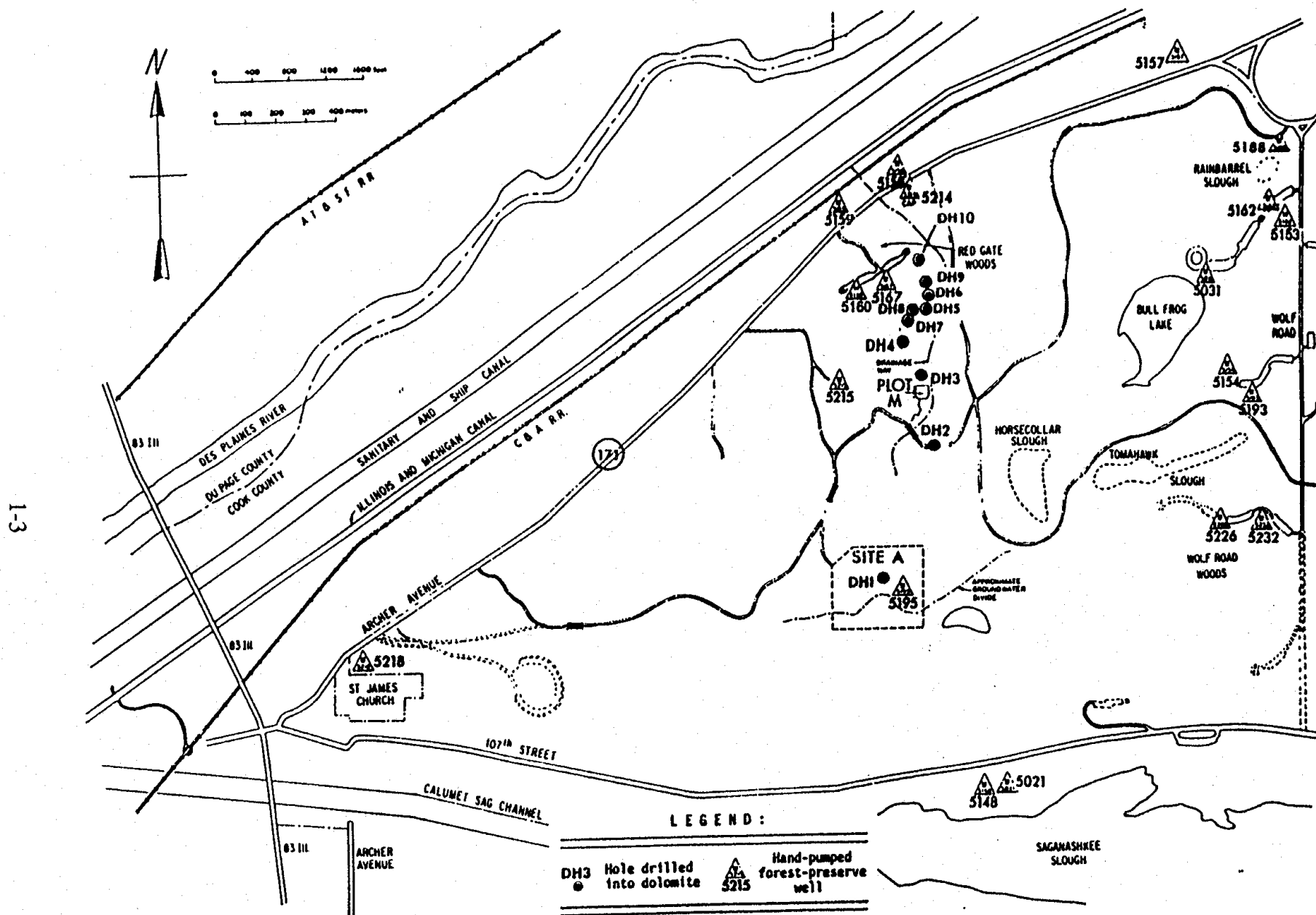


Figure 1.2 Palos Forest Preserve Showing Location of Site A/Plot M Dolomite Holes and Picnic Wells

Laboratory for disposal, but the waste buried in trenches was allowed to remain in place. Concrete sidewalls, eight feet deep, were poured around the perimeter of the burial area and a one-foot thick reinforced concrete slab was poured over the top. The concrete slab was covered with soil and seeded with grass. Both the Site A and Plot M areas were decommissioned in 1956.

In 1973, elevated levels of hydrogen-3 (as tritiated water) were detected in two nearby hand-pumped picnic wells (#5167 and #5159) and the hydrogen-3 was found to be migrating from the burial plot into the surrounding soil and aquifers. As a result, a radiological survey of the entire Palos Forest Preserve site was conducted with special emphasis on the Site A and Plot M areas.¹

The terminology used in previous reports is continued in this report. A hole drilled and completed into the glacial till is called a borehole. The soil samples obtained from the borehole are called soil cores. Some boreholes have been cased and screened to form monitoring wells. Water from such wells is called groundwater. Test wells drilled into the dolomite bedrock are called dolomite holes or deep holes. Water from such wells is called dolomite water. The hand-pumped picnic wells, which are completed into or close to the dolomite bedrock, are called water wells or picnic wells. They are identified by a location name or well number. Except for well #5160, these were in existence before this radiological and hydrological study of the area was begun.

The results of radioactivity measurements are expressed in this report in terms of picocuries per liter (pCi/L) and nanocuries per liter (nCi/L) for water samples. Radiation effective dose equivalent calculations are reported in units of millirem (mrem) or millirem per year (mrem/y). The use of the term dose throughout this report means effective dose equivalent. Other abbreviations of units are defined in the text.

1.2 Site Characteristics

Geologically, Plot M is constructed on a moraine upland which is dissected by two valleys, the Des Plaines River valley to the north and the Calumet Sag valley to the south. The upland is characterized by rolling terrain with poorly developed drainage. Streams are intermittent and drain internally or flow to one of the valleys. The area is underlain by glacial drift, dolomite, and other sedimentary rocks. The uppermost bedrock is Silurian dolomite, into which both the picnic wells and some of the monitoring wells are placed, as described in the text. The dolomite bedrock is about 200 feet thick. The overlying glacial drift has a thickness that ranges from 165 feet at Site A to zero at the Des Plaines River and Calumet Sag Canal, and some of the monitoring wells terminate in this layer. The depth to bedrock at Plot M is about 130 feet.

Hydrologically, the surface water consists of ponds and intermittent streams. When there is sufficient water, the intermittent stream that drains Plot M flows from the highest point near Site A, past Plot M, then continues near the Red Gate Woods well (Figure 1.2) and discharges into the Illinois and Michigan (I&M) Canal. The groundwater in the glacial drift and dolomite forms two distinct flow systems. The flow in the drift is controlled principally by topography. The flow in the dolomite, which is recharged by groundwater from the glacial drift, is controlled by two discharge areas, the Des Plaines River to the north and the Calumet Sag Canal to the south. Water usage in the area is confined to the hand-pumped picnic wells. These wells are open to the dolomite and are principally used in the warmer seasons.

The climate is that of the upper Mississippi valley, as moderated by Lake Michigan, and is characterized by cold winters and hot summers. Precipitation averages about 33 inches annually. The largest rainfalls occur between April and September. The average monthly temperature ranges from 21°F in January to 73°F in July. Approximately 8.4 million people reside within 50 miles of the site; the population within a five-mile radius is about 150,000. The only portion of the Palos Forest Preserve in the immediate area of Plot M and Site A that

is developed for public use is the Red Gate Woods picnic area (Figure 1.2), although small numbers of individuals use the more remote areas of the Palos Forest Preserve.

2.0 SUMMARY

The results of the ongoing environmental monitoring and surveillance program at the Palos Forest Preserve site for 2000 are presented in this report. Sample collection and analyses for radioactive substances were conducted on surface and subsurface water and for nonradioactive substances in subsurface water.

Surface water samples collected from the stream that flows around Plot M showed the same tritiated water concentration pattern seen in the past. Concentrations were at the ambient level of less than 0.1 nCi/L upstream of the Plot, increased up to 91.6 nCi/L at the seep adjacent to the Plot, then decreased to less than 0.1 nCi/L further downstream.

The tritiated water concentrations in the borehole and dolomite hole water follow a pattern consistent with that observed in the past. The tritium concentration was highest in those boreholes nearest Plot M and downgradient of the Plot. Water from eight of 16 boreholes analyzed for strontium-90 contained concentrations greater than the detection limit of 0.25 pCi/L. The elevated strontium-90 levels (up to 5.22 pCi/L) found in some boreholes are probably from the Plot, since concentrations above 0.25 pCi/L have not been observed in the groundwater due to atmospheric fallout from previous nuclear weapons testing, and no other source is known. Strontium-90 is a relatively mobile radionuclide and its presence in the borehole water is not unexpected and is probably due to migration that occurred before the Plot was capped. The strontium-90 results are consistent with those measured in the past.

Sampling of the forest preserve picnic wells shown in Figure 1.2 continued. In July 1988, the Red Gate Woods North Well (#5160) was installed as a replacement drinking water supply for the Red Gate Woods Well (#5167). The maximum and average tritium concentrations of well #5160 were 1.54 nCi/L and 1.33 nCi/L, respectively. The well opposite the entrance to Red Gate Woods (#5159) had a maximum tritium concentration of 2.20 nCi/L and an annual average concentration of 0.70 nCi/L. The other wells all averaged less than 0.1 nCi/L, except

well #5158 which averaged 0.12 nCi/L. The previous pattern of high tritium concentrations in the winter and low concentrations (less than the detection limit of 0.1 nCi/L) in the summer is not readily apparent for the wells because of the low tritium concentrations. In the calculation of annual averages, all data as measured were retained in the data base and used to compute the average.

If water equal to the Red Gate Woods North Well (#5160) average concentration of 1.33 nCi/L was the sole source of water for an individual, the annual dose from tritium would be 0.061 mrem using the DOE dose conversion factor.²¹ Consumption of one liter of this water would produce a dose of 8×10^{-5} mrem. Although the U. S. Environmental Protection Agency (EPA) drinking water regulations²² are not applicable because the picnic wells do not meet the EPA definition of a public drinking water supply, this concentration is about 7% of the EPA annual limit of 20 nCi/L. Table 4.3 provides a relative comparison of this calculated dose to natural or accepted sources of radiation.

The results of this program show that the radioactivity remaining at Site A, Plot M, and the Red Gate Woods area does not endanger the health or safety of the public visiting the site or those living in the vicinity. The potential radiation doses are very low compared to the relevant standards.

3.0 MONITORING PROGRAM

The program is designed to monitor the elevated hydrogen-3 (as tritiated water) concentrations in some of the picnic wells in the Palos Forest Preserve, determine the migration pathway of water from the burial ground to the wells, establish if other buried radionuclides or waste constituents have migrated, and characterize the radiological and non-radiological pollutant environment of the area. This is accomplished by analyzing water from wells, deep holes, boreholes, and surface streams in the area. Samples are collected from biweekly to annually, depending on past results and proximity to Plot M. During 2000, 273 samples were collected, 861 analyses were performed, and 200 field measurements were conducted. For the most part, individual results are presented in the tables and compared to control, off-site, or upstream results. Where applicable, results are compared to the U. S. Department of Energy Radiation Protection Standard of 100 mrem/y.²¹ The Site A/Plot M program follows the guidance for monitoring at DOE facilities.²³ Although it is recognized that Site A/Plot M is not a DOE facility, the same monitoring principles are applicable to this site.

The uncertainties associated with individual concentrations given in the tables are the statistical counting errors at the 95% confidence level. Because of the amount of tritium data presented on a few tables, the uncertainty values are not included. In such cases, the following uncertainties apply:

<u>Concentration (nCi/L)</u>	<u>Uncertainty (% of Conc.)</u>
0.1-1.0	40-5%
1-10	5-1%
> 10	1%

The sensitivity for the measurement of tritium in water has been improved due to new instrumentation. The current detection limit is 0.1 nCi/L.

3.1 Surface Water

Three sets of water samples were collected during 2000 from the stream that flows around Plot M, primarily during the spring when the ground was saturated. The stream was dry during scheduled sampling in the second half of the year. The sampling locations are shown in Figure 3.1. Sample Location #10 is 100 yards north of Location #9 and sample Location #11 is 200 yards north of Location #9. The outfall into the I&M Canal is approximately 600 yards north of Plot M. The samples were analyzed for tritiated water and the results are shown in Table 3.1. The same concentration pattern in the water flowing around Plot M was observed this year as in the past. Concentrations were low upstream of the Plot, increased as the stream flowed past the Plot, where it received tritiated water that leached out of the burial site, then decreased downstream due to dilution by precipitation. The outfall sample was collected prior to discharge into the I&M Canal.

Using the methodology prescribed in the DOE guidance,²¹ the committed effective dose equivalent from consumption of water can be calculated. The total quantity of an ingested radionuclide is obtained by multiplying the water concentration by the general public water ingestion rate of 730 L/y.²⁴ This annual intake is then multiplied by the 50-year Committed Effective Dose Equivalent (CEDE) factor.²⁵ The CEDE for tritium in water is 6.3×10^{-5} rem/Ci. If a hypothetical individual used water with the same tritium concentration as found in the seep (Location #6) as his sole source of water, the annual dose based on the maximum 2000 concentration of 91.6 nCi/L would be about 4.2 mrem/y and the dose based on the annual average seep concentration of 35.2 nCi/L would be 1.6 mrem/y. The DOE dose limit for the public is 100 mrem/y. Using the same calculations for concentrations at Location #9, the maximum concentration of 14.1 nCi/L would produce 0.6 mrem/y and the 2000 average concentration of 5.9 nCi/L would give a dose of 0.3 mrem/y. Consumption of one liter of water with the same annual average concentration as at Location #9 would produce a dose of 0.0004 mrem/y. In general, the tritium concentrations vary from year to year and are dependent on the amount of precipitation.



Figure 3.1 Surface Water Sampling Locations Near Plot M

TABLE 3.1

Tritiated Water Content of Stream Next to Plot M, 2000

(Concentrations in nCi/L)

Location Number*	Date Collected		
	February 23	April 20	June 13
1	< 0.1	< 0.1	< 0.1
2	< 0.1	< 0.1	< 0.1
3	< 0.1	< 0.1	9.1 ± 0.1
4	< 0.1	0.4 ± 0.1	10.5 ± 0.1
5	0.7 ± 0.1	1.1 ± 0.1	9.9 ± 0.1
6 (Seep)	9.1 ± 0.1	4.8 ± 0.1	91.6 ± 0.3
7	1.1 ± 0.1	1.3 ± 0.1	17.8 ± 0.1
8	0.5 ± 0.1	0.6 ± 0.1	4.9 ± 0.1
9	1.3 ± 0.1	1.1 ± 0.1	14.1 ± 0.1
10	1.2 ± 0.1	1.0 ± 0.1	10.5 ± 0.1
11	1.3 ± 0.1	0.9 ± 0.1	6.2 ± 0.1
Outfall**	1.0 ± 0.1	0.3 ± 0.1	< 0.1

* See Figure 3.1

** I & M Canal

The annual collection of water samples from five surface water bodies in the vicinity of Site A occurred on November 29, 2000. These are: the pond northwest of Site A; the pond southeast of Site A; Horse Collar Slough; Tomahawk Slough; and Bull Frog Lake. Most of these locations can be identified in Figure 1.2. These samples were all analyzed for tritiated water and the concentrations were all less than the detection limit of 0.1 nCi/L. The results indicate that there has been no surface migration of radioactive materials from Site A.

3.2 Subsurface Water

3.2.1 Borehole Water - Plot M

A number of the boreholes drilled in the Plot M area (Figure 3.2) were cased with plastic pipe and screens were installed (piezometers) to serve as sampling points within the glacial drift. Water samples were collected and water level measurements were made in the Plot M boreholes approximately bimonthly, weather permitting. Each borehole was emptied of water and allowed to recharge before sampling. The shallow boreholes responded to the spring precipitation as indicated by an increase in water levels followed by a drop during summer and fall when moisture was used for plant growth. The water levels in the deeper boreholes (> 100 ft) were relatively constant throughout the year. As in past years, some boreholes were occasionally dry. Borehole #11 contains a nest of piezometers at three depths; 39 feet, 68 feet, and 124 feet. The principal purpose of this arrangement is to obtain water level measurements at different depths to determine the vertical gradient of the hydraulic head.

All the water samples were analyzed for tritiated water and the results are collected in Table 3.2. The tritium concentrations varied widely as in past years. In the 40-foot deep boreholes, low tritium concentrations correlate with high water levels, apparently a dilution phenomenon. The measured water levels in the boreholes are in Table 3.3. Since the measurement of the water levels is made relative to a benchmark at the top of the well casing, a decrease in numerical value indicates a rise in water level and dilution of the tritiated water.

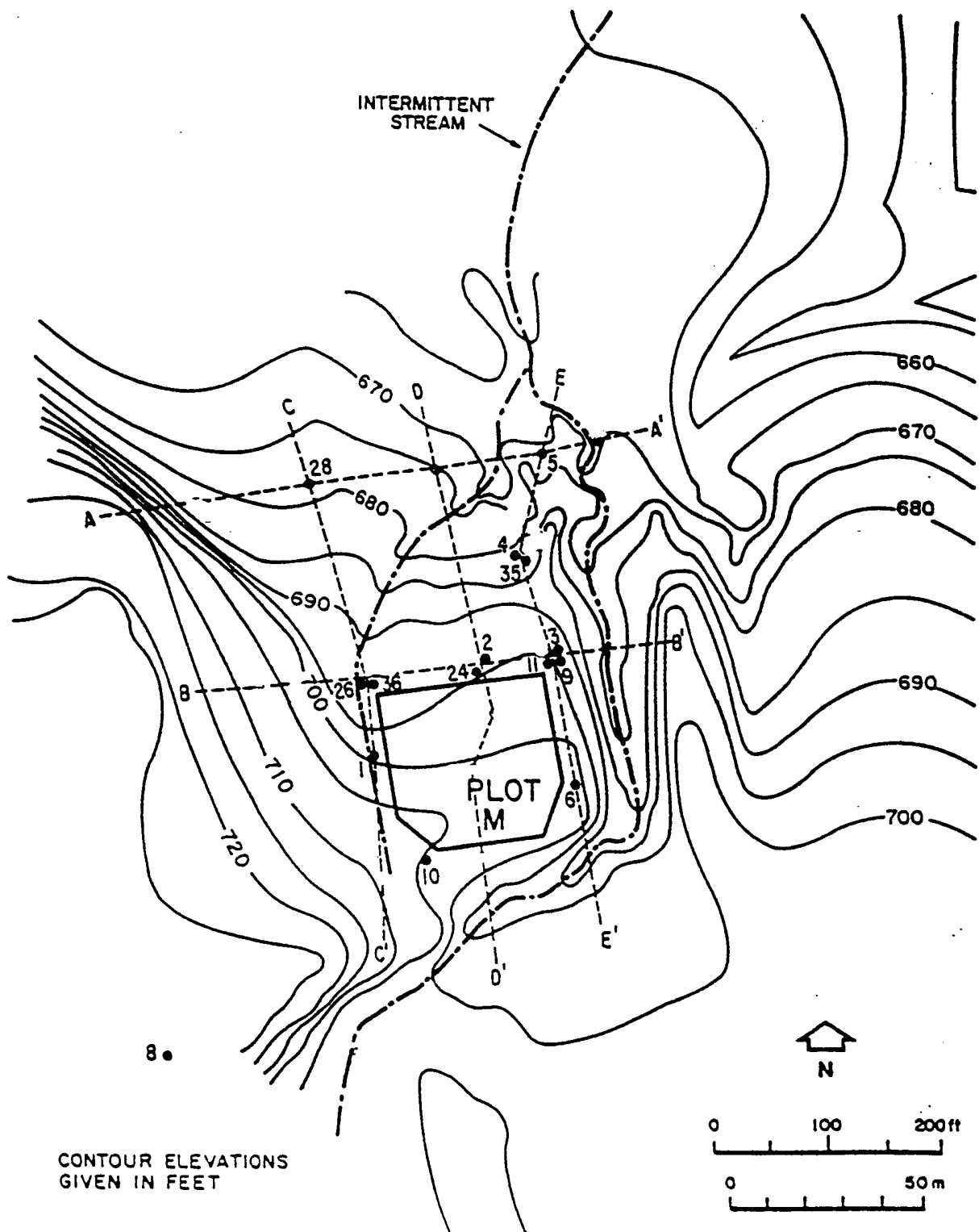


Figure 3.2 Map of Plot M Palos Site Showing Topography, Intermittent Stream, and Borehole Locations

TABLE 3.2

Tritium in Plot M Borehole Water, 2000

(Concentrations in nCi/L)

Borehole Number	Depth (ft)	Date Collected				
		February 28	April 28	June 29	August 29	December 7
1	39.46	1308.0	Dry	Dry	1305.0	Dry
2	39.41	117.7	76.4	53.9	1363.0	2280.0
3	40.00	1602.0	1439.0	1209.0	1388.0	1161.0
4	36.05	745.7	878.4	875.7	864.5	802.8
5	40.20	59.4	62.6	66.0	67.3	67.6
6	40.30	99.1	109.0	90.4	99.7	90.4
8	40.00	< 0.1	< 0.1	0.1	< 0.1	< 0.1
9	40.00*	Dry	Dry	649.8	Dry	Dry
10	40.00*	230.3	39.6	1.2	92.8	226.2
11	39.30	292.2	184.0	198.4	253.6	233.4
11	67.45	Dry	Dry	Dry	Dry	Dry
11	121.90	< 0.1	1.8	11.0	2.9	11.8
24	115.11	3.3	84.2	9.6	7.3	7.4
26	60.65	68.1	230.0	392.4	491.4	474.3
28	58.25	84.1	79.7	63.6	65.7	68.0
35	105.50	295.7	276.8	424.4	1360.0	Dry
36	120.00	10.3	5.0	6.8	7.5	11.1

* Slant hole drilled at 45° to a depth of 40 ft below the surface.

TABLE 3.3

Water Level Measurements in Boreholes Near Plot M, 2000
(Units of feet below the benchmark at the top of the well)

Borehole Number	Depth (ft)	Date Measured				
		February 28	April 28	June 29	August 29	December 7
1	39.46	39.16	Dry	Dry	38.50	Dry
2	39.41	29.00	24.55	22.91	28.14	29.05
3	40	38.97	36.41	32.24	34.55	37.90
4	36.05	24.75	18.65	14.70	19.00	21.50
5	40.20	30.13	27.50	22.89	25.00	27.30
6	40.30	38.82	34.79	31.63	33.47	36.78
8	40	35.17	35.52	35.65	37.05	35.60
11	39.30	26.39	20.86	21.14	26.60	27.75
11	67.45	Dry	Dry	Dry	Dry	Dry
11	121.90	106.15	105.70	105.95	107.90	105.75
24	115.11	80.00	80.77	81.47	88.20	98.20
26	60.65	48.10	50.27	46.61	47.90	51.05
28	58.25	57.31	56.75	55.52	55.80	55.95
35	105.50	93.78	93.37	93.48	93.45	Dry
36	120	105.41	96.93	104.98	105.49	105.47

Higher tritiated water concentrations in borehole water correlate with higher tritium concentrations in split-spoon soil cores obtained when the boreholes were constructed. In general, the magnitude of the tritium concentrations are similar to those observed over the past several years. Most of the tritium concentrations are essentially the same from sampling to sampling during the year. However, tritium concentrations in Borehole #24 and Borehole #36 continue to change significantly between sampling events. It is suspected that Borehole #24 and Borehole #36 may not be sealed properly and dilution by infiltrating precipitation may cause the tritium concentration differences. For example, when the distance from the well casing to the water level in Borehole #2 is less than 25 feet, the water levels in Borehole #24, which is next to Borehole #2, increase substantially indicating a compromise in the casing at about 25 feet below the ground surface. This is supported by the unusual water level data for Borehole #24 shown in Table 3.3.

The EPA protocols suggest that a monitoring well be purged and a sample collected within the first two hours or the sample may not be representative of the groundwater. Geological conditions at Plot M make it very difficult to follow the EPA guidance. Many of the monitoring wells at Plot M have been placed in clay-rich units with very low permeability. Recharge to these monitoring wells over a two-hour period will be insufficient to obtain an appropriate volume (up to one gallon) of water to conduct the analytical tests. The EPA criteria applies to situations where sensitive constituents such as volatile organic chemicals are of concern. Groundwater analyses at Plot M are for radioactive constituents, especially tritium which is less sensitive to chemical or physical loss from the groundwater than the volatile organic chemicals. However, the EPA sampling protocol was applied for all sampling during 2000, if operationally feasible.

As part of a search for radionuclides other than tritium in the borehole monitoring wells, sets of large volume water samples were collected to obtain greater sensitivity in the analysis. One set of samples was collected on June 29, 2000, and another set was collected December 7, 2000. Samples were collected from all boreholes that yielded sufficient water for analysis. All

samples were analyzed for strontium-90 and the results are shown in Table 3.4. Strontium-90 concentrations greater than the detection limit of 0.25 pCi/L were found in eight of the 16 sampled boreholes. Levels above 0.25 pCi/L would not be expected in this water from fallout, and no other source is known. The highest strontium-90 concentration in 2000 was found in water from Borehole #9 (5.22 pCi/L). Historically, the highest concentration was found in 1991, 10.7 pCi/L in Borehole #11 (68 feet). In the past, Borehole #6, which is between the buried waste and the stream that flows around Plot M, showed measurable strontium-90 concentrations. These data indicate small but measurable amounts of strontium-90 have migrated from the waste into the surrounding glacial drift.

3.2.2 Borehole Water - Site A

In late 1993, four boreholes (BH-41, BH-42, BH-43, and BH-44), were installed at Site A (see Figure 3.3) to improve Site A perimeter monitoring. Borehole #43 has been dry since construction. In 1994, 12 monitoring wells were constructed at Site A to support the expanded characterization of this area. With the characterization study completed in spring of 1995, the wells were transferred to the monitoring program for continued use as part of the surveillance network. These wells are also shown in Figure 3.3. Although still shown in the figure, the fence was removed in April 1998. Dedicated pumps and associated equipment were installed in July of 1995. The samples are collected semi-annually and analyzed for tritium, strontium 90, gamma-ray emitters, and metals.

The tritium results for all the Site A boreholes are collected in Table 3.5. Water levels were also measured in these boreholes and these measurements are in Table 3.6. The tritium concentrations were all low, but the pattern throughout the year was consistent. The elevated tritium levels in Borehole #41 is probably from the site landfill, while the tritium in Borehole #55 and Borehole #56 is from the CP-3 buried biological shield. The results of the strontium-90 analyses are collected in Table 3.7. The elevated strontium-90 results appear to track with elevated tritium results. For example, Boreholes #55 and #56 had measurable levels of tritium

TABLE 3.4

Strontium-90 Analyses of Borehole Water Samples Near Plot M, 2000

(Concentrations in pCi/L)

Borehole Number	Depth (ft)	Date Collected	
		June 29	December 7
1	39.46	Dry	Dry
2	39.41	0.27 ± 0.03	0.66 ± 0.03
3	40.00	< 0.25	< 0.25
4	36.05	< 0.25	< 0.25
5	40.20	< 0.25	< 0.25
6	40.30	0.51 ± 0.03	0.30 ± 0.03
8	40.00	< 0.25	< 0.25
9	40.00*	5.22 ± 0.07	Dry
10	40.00*	< 0.25	0.39 ± 0.03
11	39.30	2.11 ± 0.05	2.02 ± 0.05
11	67.45	Dry	Dry
11	121.90	< 0.25	< 0.25
24	115.11	< 0.25	0.28 ± 0.03
26	60.65	0.32 ± 0.03	< 0.25
28	58.25	< 0.25	< 0.25
35	105.50	0.49 ± 0.03	Dry
36	120.00	< 0.25	< 0.25

*Slant hole drilled at 45° to a depth of 40 ft below the surface.

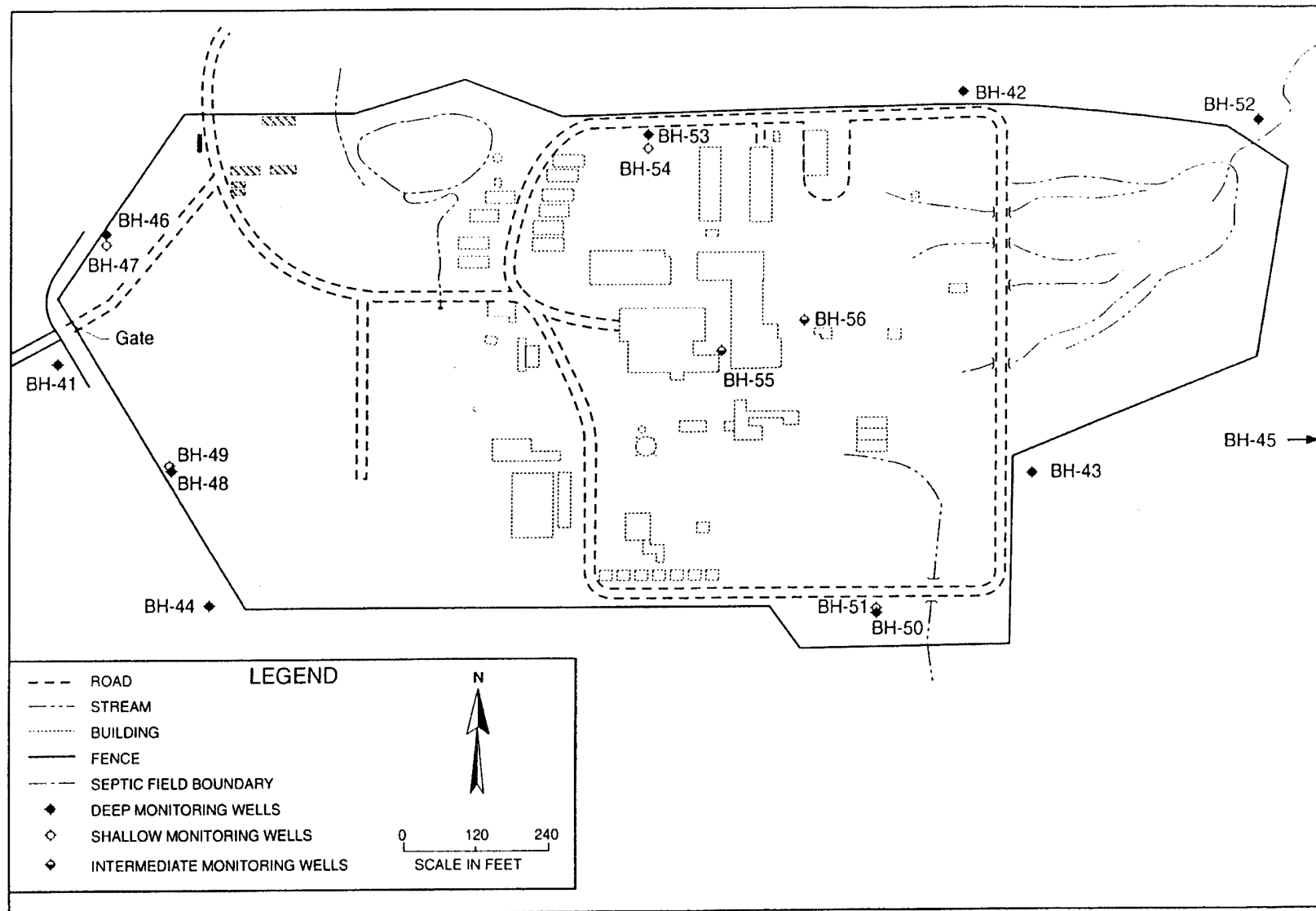


Figure 3.3 Monitoring Wells at Site A

TABLE 3.5

Tritium in Site A Borehole Water, 2000

(Concentrations in nCi/L)

Borehole Number	Depth (ft)	Date Collected	
		March 21	September 18
41	25.83	0.2	< 0.1
42	53.01	< 0.1	Dry
44	31.02	< 0.1	0.1
45	166.50	< 0.1	< 0.1
46	190.80	< 0.1	Dry
47	44.30	0.2	< 0.1
48	192.20	< 0.1	< 0.1
49	45.60	< 0.1	0.1
50	162.80	< 0.1	< 0.1
51	116.40	< 0.1	< 0.1
52	165.00	0.2	< 0.1
53	177.30	< 0.1	< 0.1
54	63.40	0.2	0.1
55	87.20	Dry	7.8
56	102.40	5.1	4.8

TABLE 3.6

Water Level Measurements in Boreholes Near Site A, 2000

(Units in feet below the benchmark at the top of the well)

Borehole Number	Depth to Bottom (ft)	Date Measured	
		March 21	September 18
41	25.83	14.74	10.55
42	53.01	51.05	Dry
43	50	Dry	Dry
44	31.02	16.01	12.75
45	166.50	143.00	143.35
46	190.80	157.36	156.98
47	44.30	19.71	16.22
48	192.20	159.79	159.28
49	45.60	22.78	15.92
50	162.80	106.86	107.25
51	116.40	101.76	102.15
52	165.00	132.23	132.45
53	177.30	150.10	150.19
54	63.40	57.18	58.19
55	87.20	Dry	79.07
56	102.40	87.75	88.06

TABLE 3.7

Strontium-90 in Site A Borehole Water Samples, 2000

(Concentrations in pCi/L)

Borehole Number*	Depth (ft)	Date Collected	
		March 21	September 18
41	25.83	0.28 ± 0.03	< 0.25
42	53.01	0.33 ± 0.03	Dry
44	31.02	< 0.25	0.31 ± 0.02
45	166.50	< 0.25	< 0.25
46	190.80	< 0.25	Dry
47	44.30	< 0.25	< 0.25
48	192.20	< 0.25	< 0.25
49	45.60	< 0.25	< 0.25
50	162.80	< 0.25	< 0.25
51	116.40	< 0.25	< 0.25
52	165.00	< 0.25	< 0.25
53	177.30	< 0.25	< 0.25
54	63.40	< 0.25	< 0.25
55	87.20	Dry	5.03 ± 0.06
56	102.40	2.01 ± 0.04	< 0.25

* See Figure 3.3.

and strontium-90 throughout the year. All gamma-ray emitters were below the detection limit of 1 pCi/L.

The Site A borehole water samples were also analyzed for total (unfiltered) metals. The results are collected in Tables 3.8 to 3.22. The concentrations of the various metals are compared to the State of Illinois Class I Ground Water Quality Standards (GWQS) ²⁶ which were selected because they represent a conservative application of the standards. Some sampling locations contained less than two results because at times the wells were dry or yielded insufficient water for the metals analysis. Those locations that contain three values include the results of a duplicate sample.

The maximum concentrations of a number of metals exceeded the GWQS in some of the monitoring wells. For a few parameters, the average concentrations also exceeded the GWQS. The exceedances are bolded in the tables. Those borehole water samples that equal or exceed the maximum GWQS are:

Antimony	Borehole #55
Arsenic	Boreholes #49, #54, and #55
Chromium	Boreholes #45 and #55
Copper	Borehole #55
Lead	Boreholes #41, #42, #45, #50, and #54
Nickel	Boreholes #45 and #55
Thallium	Borehole #55
Zinc	Boreholes #45 and #55

The average concentrations that exceeded the GWQS are:

Arsenic	Boreholes #49 and #54
Lead	Boreholes #41, #45, #54, and #55

TABLE 3.8

Chemical Constituents in Site A Borehole 41, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0030	0.0030	0.0030	0.0500
Beryllium	2	0.0009	< 0.0002	0.0017	0.0040
Cadmium	2	0.0007	0.0003	0.0011	0.0050
Chromium	2	0.0220	< 0.0000	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0130	0.0048	0.0212	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0010	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.1077	0.0483	0.1672	5.0000

TABLE 3.9

Chemical Constituents in Site A Borehole 42, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	1	-	-	< 0.0030	0.0060
Arsenic	1	-	-	0.0047	0.0500
Beryllium	1	-	-	0.0003	0.0040
Cadmium	1	-	-	0.0004	0.0050
Chromium	1	-	-	0.0440	0.1000
Copper	1	-	-	0.0341	0.6500
Lead	1	-	-	0.0090	0.0075
Mercury	1	-	-	< 0.0001	0.0020
Nickel	1	-	-	0.0502	0.1000
Selenium	1	-	-	< 0.0020	0.0500
Silver	1	-	-	< 0.0010	0.0500
Thallium	1	-	-	< 0.0015	0.0020
Zinc	1	-	-	0.1269	5.0000

TABLE 3.10

Chemical Constituents in Site A Borehole 44, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0030	0.0030	0.0030	0.0500
Beryllium	2	-	-	< 0.0001	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0020	0.0020	0.0020	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0009	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0221	0.0141	0.0301	5.0000

TABLE 3.11

Chemical Constituents in Site A Borehole 45, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0063	0.0037	0.0090	0.0500
Beryllium	2	0.0003	< 0.0001	0.0005	0.0040
Cadmium	2	0.0004	< 0.0001	0.0007	0.0050
Chromium	2	0.0903	0.0440	0.1366	0.1000
Copper	2	0.0606	0.0291	0.0921	0.6500
Lead	2	0.0129	0.0088	0.0171	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0706	0.0400	0.1012	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	0.0024	< 0.0005	0.0044	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	3.2349	0.0559	6.4140	5.0000

TABLE 3.12

Chemical Constituents in Site A Borehole 46, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	1	-	-	< 0.0030	0.0060
Arsenic	1	-	-	0.0030	0.0500
Beryllium	1	-	-	< 0.0001	0.0040
Cadmium	1	-	-	< 0.0001	0.0050
Chromium	1	-	-	0.0440	0.1000
Copper	1	-	-	< 0.0170	0.6500
Lead	1	-	-	0.0020	0.0075
Mercury	1	-	-	< 0.0001	0.0020
Nickel	1	-	-	0.0400	0.1000
Selenium	1	-	-	< 0.0020	0.0500
Silver	1	-	-	< 0.0010	0.0500
Thallium	1	-	-	< 0.0015	0.0020
Zinc	1	-	-	0.0519	5.0000

TABLE 3.13

Chemical Constituents in Site A Borehole 47, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	3	-	-	< 0.0030	0.0060
Arsenic	3	0.0030	0.0030	0.0030	0.0500
Beryllium	3	-	-	< 0.0001	0.0040
Cadmium	3	-	-	< 0.0001	0.0050
Chromium	3	0.0440	0.0440	0.0440	0.1000
Copper	3	-	-	< 0.0170	0.6500
Lead	3	0.0023	0.0020	0.0025	0.0075
Mercury	3	-	-	< 0.0001	0.0020
Nickel	3	0.0400	0.0400	0.0400	0.1000
Selenium	3	-	-	< 0.0020	0.0500
Silver	3	0.0012	< 0.0005	0.0017	0.0500
Thallium	3	-	-	< 0.0015	0.0020
Zinc	3	0.0301	0.0168	0.0421	5.0000

TABLE 3.14

Chemical Constituents in Site A Borehole 48, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0030	0.0030	0.0030	0.0500
Beryllium	2	-	-	< 0.0001	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0020	0.0020	0.0020	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0006	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0202	0.0110	0.0294	5.0000

TABLE 3.15

Chemical Constituents in Site A Borehole 49, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	3	-	-	< 0.0030	0.0060
Arsenic	3	0.0645	0.0030	0.1033	0.0500
Beryllium	3	-	-	< 0.0001	0.0040
Cadmium	3	-	-	< 0.0001	0.0050
Chromium	3	0.0440	0.0440	0.0440	0.1000
Copper	3	-	-	< 0.0170	0.6500
Lead	3	0.0020	0.0020	0.0020	0.0075
Mercury	3	-	-	< 0.0001	0.0020
Nickel	3	0.0400	0.0400	0.0400	0.1000
Selenium	3	-	-	< 0.0020	0.0500
Silver	3	-	-	< 0.0007	0.0500
Thallium	3	-	-	< 0.0015	0.0020
Zinc	3	0.0155	0.0110	0.0242	5.0000

TABLE 3.16

Chemical Constituents in Site A Borehole 50, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0032	0.0030	0.0035	0.0500
Beryllium	2	-	-	< 0.0002	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0063	0.0048	0.0078	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0005	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0346	0.0331	0.0362	5.0000

TABLE 3.17

Chemical Constituents in Site A Borehole 51, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0092	0.0090	0.0094	0.0500
Beryllium	2	-	-	< 0.0001	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0020	0.0020	0.0020	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0006	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0135	0.0110	0.0160	5.0000

TABLE 3.18

Chemical Constituents in Site A Borehole 52, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0030	0.0030	0.0030	0.0500
Beryllium	2	-	-	< 0.0001	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0020	0.0020	0.0020	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0007	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0110	0.0110	0.0110	5.0000

TABLE 3.19

Chemical Constituents in Site A Borehole 53, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0096	0.0085	0.0107	0.0500
Beryllium	2	-	-	< 0.0001	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0020	0.0020	0.0020	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0008	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0144	0.0128	0.0161	5.0000

TABLE 3.20

Chemical Constituents in Site A Borehole 54, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0076	0.0030	0.0123	0.0500
Beryllium	2	-	-	< 0.0002	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0049	0.0020	0.0079	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	-	-	< 0.0007	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0567	0.0208	0.0925	5.0000

TABLE 3.21

Chemical Constituents in Site A Borehole 55, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	1	-	-	0.0150	0.0060
Arsenic	1	-	-	0.0030	0.0500
Beryllium	1	-	-	< 0.0001	0.0040
Cadmium	1	-	-	0.0002	0.0050
Chromium	1	-	-	0.3965	0.1000
Copper	1	-	-	1.1420	0.6500
Lead	1	-	-	0.0040	0.0075
Mercury	1	-	-	< 0.0001	0.0020
Nickel	1	-	-	0.7657	0.1000
Selenium	1	-	-	< 0.0020	0.0500
Silver	1	-	-	< 0.0005	0.0500
Thallium	1	-	-	0.0030	0.0020
Zinc	1	-	-	2.7460	5.0000

TABLE 3.22

Chemical Constituents in Site A Borehole 56, 2000

(Concentrations in mg/L)

Constituent	No. of Samples	Concentrations			GWQS
		Avg.	Min.	Max.	
Antimony	2	-	-	< 0.0030	0.0060
Arsenic	2	0.0035	0.0030	0.0041	0.0500
Beryllium	2	-	-	< 0.0001	0.0040
Cadmium	2	-	-	< 0.0001	0.0050
Chromium	2	0.0440	0.0440	0.0440	0.1000
Copper	2	-	-	< 0.0170	0.6500
Lead	2	0.0020	0.0020	0.0020	0.0075
Mercury	2	-	-	< 0.0001	0.0020
Nickel	2	0.0400	0.0400	0.0400	0.1000
Selenium	2	-	-	< 0.0020	0.0500
Silver	2	0.0099	0.0048	0.0150	0.0500
Thallium	2	-	-	< 0.0015	0.0020
Zinc	2	0.0135	0.0135	0.0135	5.0000

Borehole #54 is a shallow monitoring well near the former vehicle services buildings, while Borehole #55 monitored the buried CP-3 biological shield where building debris and other material were buried. Borehole #45 was installed as an upgradient location, but has the most exceedances. Over the past two years, the samples from this well have had greater turbidity than previous years and increased metals concentrations are frequently associated with increased turbidity. It is thought that sample turbidity in this and other boreholes may be contributing to the exceedances of the groundwater quality standards.

3.2.3 Dolomite Hole Water

At the present time, 15 wells are cased into the dolomite zone to monitor the movement of any radionuclides in this aquifer. Most of the dolomite holes are located north of Plot M and east of the Red Gate Woods North Well (#5160), as shown in Figure 1.2 and/or Figure 3.4. Water was normally collected from the dolomite holes quarterly. All samples were analyzed for tritiated water and the results are in Table 3.23. Water levels were also measured in the dolomite holes and these measurements are in Table 3.24. Due to transition of staff, the third quarter samples were inadvertently not collected and the second quarter water level measurements lost. Some of the fourth quarter samples could not be collected because of heavy snow in mid-December which prevented access to some part of the site.

In December 1993, DH 11 through DH 14 were grouted with cement to 535 ft MSL. The purpose of the grouting was to seal off the lower dolomite fractures that are connected to the hand-pumped picnic wells in the Red Gate Woods area. It is speculated that these lower fractures allow elevated levels of tritiated water in the dolomite holes to migrate to the picnic well. In addition, covers to three of the dolomite holes were repaired.

In September 1994, three of the dolomite holes, DH 6, DH 7, and DH 8, were abandoned because the wells were not sealed into the dolomite, but were open to the glacial till. As such, they were not sampling dolomite water. Borehole #40 was also abandoned at the same time

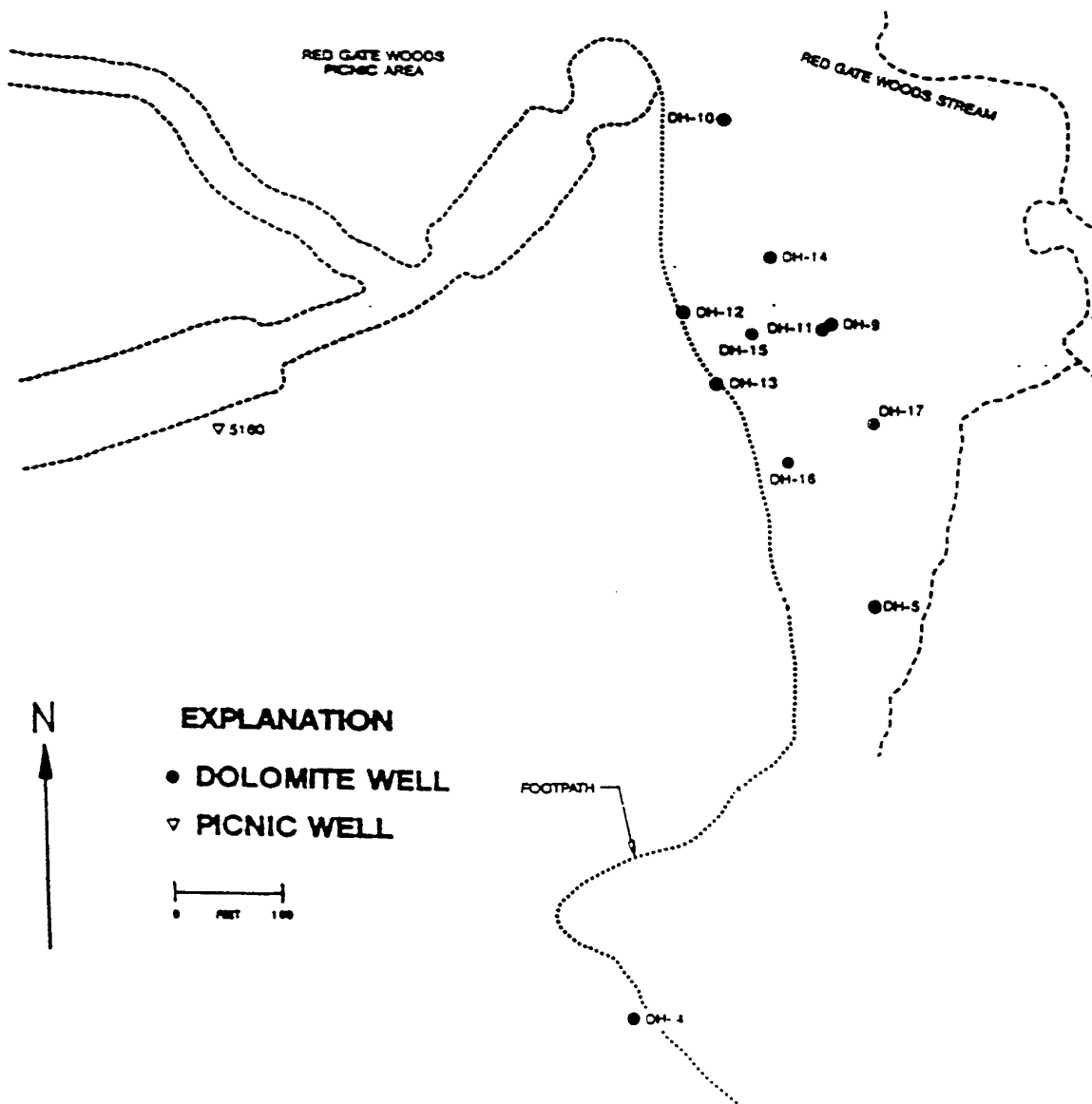


Figure 3.4 Locations of Dolomite Holes North of Plot M

TABLE 3.23

Tritium in Site A Dolomite Holes, 2000

(Concentrations in nCi/L)

Dolomite Hole Number	Date Collected		
	March 16	June 22	December 19
1	< 0.1	< 0.1	Snow*
2	< 0.1	< 0.1	Snow*
3	1.7	1.6	Snow*
4	0.1	< 0.1	Snow*
5	< 0.1	< 0.1	< 0.1
9	0.4	0.2	0.8
10	2.2	2.2	1.1
11	2.4	2.3	2.4
12	2.7	2.8	2.6
13	2.0	2.0	1.9
14	3.1	3.0	3.0
15	1.9	1.8	1.8
17	0.4	0.4	0.3
18	< 0.1	< 0.1	Snow*

*Locations inaccessible due to heavy snow.

TABLE 3.24

Water Level Measurements in Dolomite Holes 2000

(Units in feet below the benchmark at the top of the well)

Dolomite Hole Number	Date Measured	
	March 16	December 19
1	162.05	Snow*
2	140.50	Snow*
3	99.01	Snow*
4	94.30	Snow*
5	79.06	79.50
9	73.09	73.16
10	64.81	65.33
11	76.95	77.50
12	78.02	78.50
13	78.64	79.17
14	73.02	73.50
15	80.63	81.15
16	Dry	Dry
17	75.90	76.40
18	144.32	Snow*

*Locations inaccessible due to heavy snow.

because the well casing was probably compromised. In accordance with well sealing procedures established by the Illinois Environmental Protection Agency, the casing was filled with clean gravel and sand, sealed with bentonite, the casing removed to three feet below the ground level, and backfilled with clean soil.

The results of the tritiated water analyses of the dolomite holes are consistent with concentrations measured in the past. Nine of the dolomite holes had elevated tritiated water concentrations. The highest tritium levels are in the eight dolomite holes, DH 9 to DH 15 and DH 17, which are the furthest north and near the surface stream that flows next to Plot M (see Section 3.2). The distribution of tritium in these wells supports the USGS interpretation²⁷ that a large tritium plume underlies the stream. The plume has spread downward as well as downgradient resulting in the current configuration of the tritium concentrations in the dolomite. The other dolomite hole with elevated tritium is DH 3, which is immediately downgradient from Plot M. Previous analyses of soil core samples indicated the presence of tritium down to the drift-dolomite interface at DH 3.

Past sampling practices indicate different tritium results are obtained depending on sampling methods. Within the dolomite hole wells north of Plot M, collection of the sample near the top of the water column (shallow) will result in measurable tritium concentrations. Sampling at the bottom of the water column (deep) will result in very low or non-detectable tritium concentrations. In addition, samples collected at any depth after purging, removing up to 700 L of water before sampling, will result in samples with tritium concentrations that are very low or non-detectable.

The explanation for the differences in tritium concentration is that the tritium is moving horizontally in the uppermost fractures of the dolomite. The tritium is in a narrow band of the water column in the dolomite hole. Where a sample is collected with a bailer, this tritium zone is the area sampled resulting in measurable levels of tritium in the sample. However, when the dolomite hole is purged, the volume of recharge water from other fractures that are free of

tritium is such that the tritium concentration in the sample is generally at the detection limit. If the next quarter sample is collected without purging, the tritium concentration is at the same level as earlier samples indicating that the fracture flow rapidly reached an equilibrium condition.

3.2.4 Well Water

In July 1988, a well was installed in the Red Gate Woods picnic area (#5160) to replace the existing well (#5167) as a drinking water supply for visitors to this area. This is referred to as the Red Gate Woods North Well (#5160). This well was to be cased 20 feet into the dolomite to seal off fractures assumed to contain tritiated water. Previous experiments had indicated that the tritium moved at the drift-dolomite interface and in the uppermost fractures in the dolomite. Extending the casing 20 feet into the dolomite and cementing the void between the casing and dolomite was expected to prevent tritium from entering the new well via these fractures. But this was not successful. In addition, placement of a new pump mechanism on the new well was expected to eliminate the elevated lead concentrations found occasionally in the old well that resulted from corrosion of the old pump mechanism. The pump mechanism was removed from the old Red Gate Woods well to prevent its use by the public; however, the well was maintained as a sampling location for this monitoring program. In addition, the lower portion of the Red Gate Woods Well (#5167) was filled with grout to seal the lower fractures and prevent communication of water with the new well. Since 1989, iron, lead, manganese, and zinc have regularly exceeded the GWQS at the Red Gate Woods North Well (#5160) in addition to tritium.

Sampling was conducted throughout the year at the forest preserve picnic wells located north of Plot M and shown in Figure 1.2. All the samples were analyzed for tritiated water and the results are listed in Table 3.25. Occasionally, the pumps were not operable (broken) and therefore no samples were collected. Well #5215 (Guard Post) became inoperative early in 1999, was not repairable, and was removed from the monitoring program. In addition, the Red

TABLE 3.25

Tritiated Water Content of Wells Near Site A/Plot M, 2000

(Concentrations in nCi/L)

Date Collected	Red Gate North #5160	Opposite Red Gate #5159	300 yds. East Red Gate #5158
January 5	1.18	2.20	0.16
January 20	1.22	1.75	-
February 2	1.21	Broken	< 0.1
February 16	1.15	1.56	-
March 1	1.25	1.50	0.14
March 16	1.17	1.21	-
April 6	1.19	1.03	0.15
April 20	1.09	1.06	-
May 4	1.30	0.41	0.11
May 17	1.34	0.54	-
June 7	1.39	0.54	0.13
June 23	Broken	0.41	-
July 5	Broken	0.39	0.12
July 19	1.40	0.35	-
August 7	1.40	0.33	< 0.1
August 16	1.41	0.34	-
September 6	1.49	0.30	0.12
September 20	1.45	0.28	-
October 4	1.43	0.30	0.11
October 18	1.54	0.28	-
November 1	1.42	0.23	< 0.1
November 16	1.46	0.20	-
December 6	1.50	0.23	0.12
December 20	Snow	Snow	-
Average	1.33	0.70	0.12

Gate Woods North Well (#5160) was not available to the public most of 1999 and 2000 because of high fecal coliform levels. The tritium concentrations in the wells have decreased to the level where the earlier pattern of high concentrations in the winter and low concentrations in the summer is not readily detectable. The maximum and average tritium concentrations since 1976 for wells #5160, #5167, and #5159 are presented in Table 3.26. The tritium concentration over the past few years is illustrated in Figure 3.5, which is a plot of the tritium concentrations in wells #5160, #5167, and #5159 for the past nine years. The tritium concentration in the Red Gate Woods North Well (#5160) increased to about 2.2 nCi/L in November 1995 and has shown a gradual decrease in concentration during 1996, 1997, and 1998, remained constant throughout all of 1999, but gradually increased in 2000. This is contrary to past concentration behavior of higher in the winter and lower in the summer. The concentration pattern acts as if there were a localized reservoir of tritium. Before the Red Gate Woods Well (#5167) was sealed, the tritium concentrations had decreased to below the detection limit. The tritium concentrations in the well opposite Red Gate Woods (#5159) are more irregular and may be related to the amount of precipitation. The final result in 1999 is significantly higher than the rest of the year, but decreased consistently through 2000. The annual average tritium concentrations are slightly higher in the new well (#5160) compared to the old well (#5167), see Table 3.26.

The other wells, although also downgradient from Plot M, are evidently too far from the Plot to show measurable tritium concentrations. Two of the picnic wells, #5159 and #5158, were not available to the public during the year. The Forest Preserve District of Cook County had removed the pump handles due to high fecal coliform concentrations in the well water. A set of picnic well water samples was collected on May 25, 2000, and another set on November 28, 2000, from the wells on the east and south sides of the Palos Forest Preserve. The sampled wells were #5021, #5031, #5149, #5153, #5154, #5162, #5188, #5193, #5226, and #5232 in Figure 1.2. All the tritium results were less than the detection limit of 0.1 nCi/L.

TABLE 3.26

Annual Maximum and Average Tritium Concentrations
in the Red Gate Woods Wells

(Concentrations in nCi/L)

Year	Red Gate Woods North (#5160)*		Red Gate Woods (#5167)		Opposite Red Gate Woods (#5159)	
	Maximum	Annual Average	Maximum	Annual Average	Maximum	Annual Average
1976			11.0	6.1	4.4	1.90
1977			9.2	8.2	5.1	2.10
1978			12.2	7.5	2.4	1.30
1979			11.4	7.1	2.6	0.96
1980			8.8	7.0	1.6	1.02
1981			9.6	4.3	2.1	1.06
1982			11.0	4.8	3.2	0.79
1983			5.1	2.2	1.8	0.95
1984			2.7	1.2	1.70	0.70
1985			2.5	1.0	1.60	0.69
1986			3.4	1.3	0.94	0.60
1987			3.3	1.6	0.73	0.54
1988	0.30	0.19	0.48	0.18	1.91	1.32
1989	0.41	0.27	0.46	0.30	0.92	0.61
1990	0.52	0.22	0.25	0.12	2.07	1.14
1991	0.80	0.35	0.21	< 0.1	2.26	1.43
1992	0.54	0.25	0.13	< 0.1	2.29	1.15
1993	0.57	0.23	< 0.1	< 0.1	1.63	0.87
1994	0.40	0.31	< 0.1	< 0.1	0.46	0.30
1995	2.26	0.51	< 0.1	< 0.1	0.54	0.40
1996	2.19	1.56	Closed/Sealed November 1995		0.55	0.33
1997	1.26	1.00			1.13	0.35
1998	1.23	1.03			0.72	0.47
1999	1.22	1.07			2.14	0.45
2000	1.54	1.33			2.20	0.70

*The replacement well (#5160) was installed in July 1988.

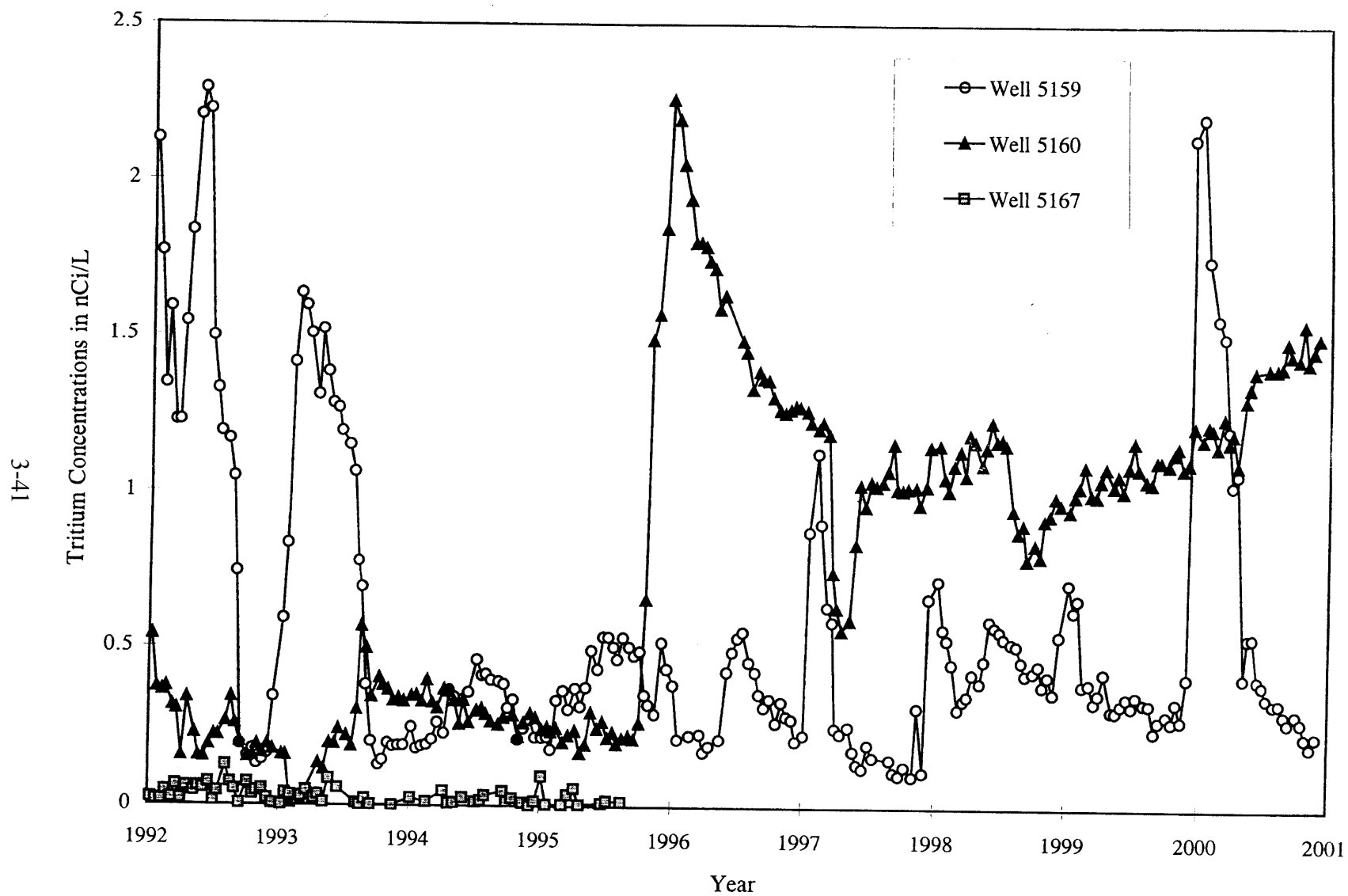


Figure 3.5 Tritiated Water Concentrations in Red Gate Woods (#5167), Opposite Red Gate Woods (#5159), and Red Gate Woods North (#5160) Wells From 1992 Through 2000

The Red Gate Woods North Well (#5160) is the location which provided the greatest potential radiation exposure to the public in 2000. If water equal to the Red Gate Woods North well average concentration of 1.33 nCi/L were the sole source of water for an individual, the annual dose from the tritium would be 0.061 mrem. If an individual consumed one liter of this water, the dose would be 8×10^{-5} mrem.

Quarterly water samples collected from the picnic wells were also analyzed for isotopic uranium. The uranium results are presented in Table 3.27. The range of concentrations is similar to that found previously and is consistent with concentrations of uranium found in well water away from Site A/Plot M.

Samples were collected quarterly from the Red Gate Woods North Well (#5160) were analyzed unfiltered for a number of inorganic constituents. The results are found in Table 3.28. Because this picnic well is used as a drinking water supply, the limits used were the State of Illinois concentrations of chemical constituents in drinking water.²⁶ The constituents in Table 3.28 that do not have a specific limit are provided for completeness. Elevated levels of copper, iron, lead, and zinc were found in three of the four sampling events, but their presence may be related to the decomposition of the well pump materials. There is no evidence that the source of the elevated inorganic constituents is from Plot M, since these metals are not found in the deepholes between Plot M and the Red Gate Woods wells. Although the limits are exceeded for manganese, this is due to natural levels. These concentrations are not unusual for dolomite water in northeast Illinois.²⁸

Four wells in private homes near 107th and Archer Avenue were sampled on September 19, 2000, and analyzed for gross alpha, gross beta, tritium, and gamma-ray-emitting radionuclides. These wells are in the same dolomite zone as the picnic wells. The gross alpha and gross beta activities were in the normal range of naturally-occurring radionuclides in groundwater for this part of Illinois. The gross alpha activities ranged from 2.6 to 3.3 pCi/L and the gross beta activities ranged from 9.0 to 13.0 pCi/L. All the tritium results were less

TABLE 3.27

Uranium Content of Well Water Near Plot M, 2000

(Concentrations in pCi/L)

Well Number*	Location	Date Collected			
		February 2	May 4	August 7	November 1
5160	Red Gate Woods North	0.01 ± 0.04	0.04 ± 0.04	0.03 ± 0.04	0.04 ± 0.06
5159	Opposite Entrance to Red Gate Woods	Broken	0.73 ± 0.11	0.10 ± 0.08	0.04 ± 0.06
5158	300 yds NE of Entrance to Red Gate Woods	-	0.04 ± 0.04	-	-

* See Figure 1.2

TABLE 3.28

Inorganic Constituents in Red Gate Woods North Well Water (#5160), 2000

(Concentrations in mg/L)

Inorganic Constituent	February	May	August	November	GWQS*
Arsenic	< 0.00030	< 0.0030	< 0.0030	< 0.0030	0.05
Barium	0.026	< 0.018	< 0.018	< 0.018	2.0
Beryllium	< 0.0001	< 0.0001	< 0.0001	< 0.0001	0.004
Cadmium	0.0014	0.0035	0.0014	< 0.0001	0.005
Chloride	46	37	31	31	200
Chromium	< 0.044	< 0.044	< 0.044	< 0.044	0.1
Cobalt	< 0.026	< 0.026	< 0.026	< 0.026	1.0
Copper	0.023	1.50	7.62	< 0.017	0.65
Fluoride	0.242	0.134	0.104	0.080	4.0
Iron	50.4	101.0	24.9	0.06	5.0
Lead	0.135	0.344	0.132	< 0.002	0.0075
Manganese	0.448	0.683	1.360	< 0.017	0.15
Mercury	< 0.0001	< 0.0001	< 0.0001	< 0.0001	0.002
Nickel	< 0.04	< 0.04	0.018	< 0.04	0.1
pH	7.39	8.21	8.02	8.32	6.5-9.0
Silver	0.0008	0.0030	0.0047	< 0.0005	0.05
Sulfate	54	61	157	130	400
Thallium	< 0.0015	< 0.0015	< 0.0015	< 0.0015	0.002
Vanadium	< 0.024	0.024	0.062	< 0.024	-
Zinc	24.1	131.3	67.9	0.076	5.0

* State of Illinois Class I Ground Water Quality Standards (GWQS).²⁶

than the detection limit of 0.1 nCi/L and gamma-ray spectrometric analysis of the water did not detect any radionuclides associated with activities at Site A/Plot M above the detection limit of 1 pCi/L. These results indicate that no radioactivity used or generated at Site A/Plot M has entered this water system.

4.0 SUMMARY OF POTENTIAL RADIATION DOSE AND RISK ESTIMATES

4.1 Dose Estimates

The dose from drinking water to an individual exposed to radionuclides associated with Plot M can be estimated employing the DOE methodology. If a hypothetical individual were exposed continuously to tritium at various locations near Plot M, the dose could be estimated. Assuming a person drank water from Location #9 or the seep (Location #6), or drank water from well #5160, the dose from exposure for all of 2000 at the maximum and annual average concentrations is collected in Table 4.1. This scenario assumes that the individual's sole source of water is at the identified location.

A more meaningful estimation is for the occasional visitor to the Plot M area. Assuming a visitor drinks one liter of water from the surface stream or picnic wells, the dose from this exposure is estimated and presented in Table 4.2. The maximum total dose received by an occasional visitor is the combination of surface water and drinking water from the Red Gate Woods North Well (#5160). This maximum dose would be 0.0001 mrem per visit.

In order to put the doses into perspective, comparisons can be made to annual average doses received by the public from natural or accepted sources of radiation. These are listed in Table 4.3. It is obvious that the magnitude of the doses potentially received near Plot M from residual radioactive substances remaining from work conducted in this area are insignificant compared to these sources.

4.2 Risk Estimates

Risk estimates of possible health effects from radiation doses to the public from Plot M have been made to provide another perspective in interpreting the radiation doses. Estimated for carcinogenic risk, the risk of contracting cancer from these exposures, is included in

TABLE 4.1

Dose From Continuous Exposure to Tritium at Selected Locations, 2000

Pathway	Maximum				Annual Average				Average Carcinogenic Risk
	Conc.		Dose		Conc.		Dose		
Surface Water									
Seep	91.6	nCi/L	4.2	mrem/y	35.2	nCi/L	1.6	mrem/y	1 x 10 ⁻⁶
Location #9	14.1	nCi/L	0.6	mrem/y	5.9	nCi/L	0.3	mrem/y	2 x 10 ⁻⁷
Well Water									
Red Gate Woods North (#5160)	1.54	nCi/L	0.071	mrem/y	1.33	nCi/L	0.061	mrem/y	4 x 10 ⁻⁸

TABLE 4.2

Estimates of Tritium Exposures to a Casual Visitor to Plot M, 2000

Pathway	Quantity	Maximum Dose	Annual Average	Average Carcinogenic Risk
Surface Water				
Seep	One Liter	0.006 mrem	0.002 mrem	2×10^{-9}
Location #	One Liter	0.0008 mrem	0.0004 mrem	3×10^{-10}
Well Water				
Red Gate Woods North (#5160)	One Liter	0.00010 mrem	0.00008 mrem	5×10^{-11}

TABLE 4.3

Annual Average Dose Equivalent in the U. S. Population*	
Source	(mrem)
Natural Sources	
Radon	200
Internal (^{40}K and ^{226}Ra)	39
Cosmic	28
Terrestrial	28
Medical	
Diagnostic X-rays	39
Nuclear Medicine	14
Consumer Products	
Domestic Water Supplies, Building Materials, etc.	10
Occupational (Medical Radiology, Industrial Radiography, Research, etc.)	1
Nuclear Fuel Cycle	< 1
Fallout	< 1
Other Miscellaneous sources	< 1
Total	360

*NCRP report No. 93.²⁹

Table 4.1 and Table 4.2 for the average exposure scenario. Based on the BIER V report,³⁰ a dose of one mrem/y equates to an increased risk of 0.7×10^{-6} . This conversion ratio is used in these tables. The risks are estimated to be in addition to the normal incident rate of cancer in the general population. For example, a carcinogenic risk of 10^{-7} would mean one additional cancer to 10,000,000 people exposed under the prescribed conditions. The EPA environmental protection standards are generally based on an acceptable risk between 10^{-4} and 10^{-6} . This would imply that a risk of greater than 10^{-4} would be unacceptable and a risk of less than 10^{-6} would be acceptable. Examination of Table 4.1 indicates that even under the very conservative assumptions of sole source use of the water at Plot M annual average concentrations, the risk is less than the EPA recommendation. For the Table 4.2 hypothetical dose to an occasional visitor of 0.00001 mrem, the risk would be about 10^{-11} . The risk from exposure to radionuclides at Plot M can be compared to the risk associated with natural events. A few examples are collected in Table 4.4. The risk from the naturally-occurring sources of radioactivity listed in Table 4.3 is estimated to be about one additional cancer in a population of 8,000. Therefore, the monitoring program results have established that radioactivity at Plot M is very low and does not endanger the health or safety of those living in the area or visiting the site.

TABLE 4.4

Risk of Death From Natural Events	
Cause	Risk
Lightning Strike	5×10^{-8}
Tornado	1×10^{-7}
Flood	1×10^{-7}
Hurricane	2.5×10^{-7}
Drowning	8×10^{-6}
Air Travel	3×10^{-6}
Firearms	2×10^{-6}

5.0 REFERENCES

1. Golchert, N. W. and Sedlet, J., *Formerly Utilized MED/AEC Sites Remedial Action Program - Radiological Survey of Site A, Palos Park Forest Preserve, Chicago, Illinois*, U. S. Department of Energy Report DOE/EV-0005/7 (April 1978).
2. Golchert, N. W., Sedlet, J., and Hayes, K. A., *Environmental Surveillance of the Palos Park Forest Preserve*, U. S. Department of Energy Report ANL-83-6 (January 1983).
3. Golchert, N. W. and Sedlet, J., *Site Surveillance and Maintenance Program for Palos Park - Report for 1982*, Argonne National Laboratory (available from the authors) (April 1984).
4. Golchert, N. W. and Sedlet, J., *Site Surveillance and Maintenance Program for Palos Park - Report for 1983*, Argonne National Laboratory (available from the authors) (June 1984).
5. Golchert, N. W. and Sedlet, J., *Site Surveillance and Maintenance Program for Palos Park - Report for 1984*, Argonne National Laboratory (available from the authors) (April 1985).
6. Golchert, N. W. and Sedlet, J., *Site Surveillance and Maintenance Program for Palos Park - Report for 1985*, U. S. Department of Energy Report ANL-86-25 (April 1986).
7. Golchert, N. W., *Site Surveillance and Maintenance Program for Palos Park - Report for 1986*, U. S. Department of Energy Report ANL-87-8 (April 1987).
8. Golchert, N. W., *Site Surveillance and Maintenance Program for Palos Park - Report for 1987*, U. S. Department of Energy Report ANL-88-12 (April 1988).

9. Golchert, N. W., *Site Surveillance and Maintenance Program for Palos Park - Report for 1988*, U. S. Department of Energy Report ANL-89/7 (April 1989).
10. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1989*, U. S. Department of Energy Report ANL-90/7 (April 1990).
11. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1990*, U. S. Department of Energy Report ANL-91/2 (May 1991).
12. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1991*, U. S. Department of Energy Report ANL-92/13 (May 1992).
13. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1992*, U. S. Department of Energy Report ANL-93/4 (May 1993).
14. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1993*, U. S. Department of Energy Report ANL-94/9 (May 1994).
15. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1994*, U. S. Department of Energy Report ANL-95/7 (May 1995).
16. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1995*, U. S. Department of Energy Report ANL-96/2 (June 1996).
17. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1996*, U. S. Department of Energy Report ANL-97/5 (May 1997).
18. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1997*, U. S. Department of Energy Report ANL-98/1 (May 1998).

19. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1998*, U. S. Department of Energy Report, ANL-99/2 (May 1999).
20. Golchert, N. W., *Surveillance of Site A and Plot M - Report for 1999*, U. S. Department of Energy Report, ANL-00/3 (May 2000).
21. U. S. Department of Energy, "Radiation Protection of the Public and the Environment," *DOE Order 5400.5*, February 8, 1990.
22. U. S. Environmental Protection Agency, "National Primary Drinking Water Regulations," 40 CFR Part 141.
23. U. S. Department of Energy, "Environmental Regulatory Guide for Radiological Effluent Monitoring and Environmental Surveillance," *DOE/EH-0173T*, January 1991.
24. International Commission on Radiological Protection, "Reference Man: Anatomical, Physiological, and Metabolic Characteristics," *ICRP Publication 23*, Pergamon Press, New York, NY (1975).
25. U. S. Department of Energy, "Internal Dose Conversion Factors for Calculation of Dose to the Public," *DOE/EH-0071*, July 1988.
26. Illinois Pollution Control Board, Illinois Administrative Code, Title 35, Subtitle F; Chapter I, Part 620, "Ground Water Quality," September 15, 1994.
27. Nicholas, J. R. and Healy, R. W., "Tritium Migration From a Low-Level Radioactive-Waste Disposal Site Near Chicago, Illinois," U. S. Geological Survey Water-Supply Paper 2333, 1988.

28. Final report - Tritium Migration and Hydrogeological Studies in the Vicinity of Plot M - Palos Forest Preserve, International Technology Corporation Report, Project No. 313433, August 1994.
29. National Council on Radiation Protection and Measurements, *Ionizing Radiation Exposure of the Population of the United States*, NCRP Report No. 93, September 1, 1987.
30. Committee on Biological Effects of Ionizing Radiation, *Health Effects on Populations of Exposure to Low Levels of Ionizing Radiation - BEIR V Report*, National Academy Press, Washington, 1990.
31. Golchert, N. W. and Kolzow, R. G., *Argonne National Laboratory-East Site Environmental Report for Calendar Year 1999*, U. S. Department of Energy Report ANL-00/2 (September 2000).

6.0 APPENDICES

6.1 Quality Assurance Program

All nuclear instrumentation is calibrated with standardized sources obtained from or traceable to the U. S. National Institute of Standards and Technology (NIST). The equipment is checked prior to the sample measurements with secondary counting standards to insure proper operation. Samples were periodically analyzed in duplicate or with the addition of known amounts of a radionuclide to check precision and accuracy. Intercomparison samples distributed by the DOE Environmental Measurements Laboratory Quality Assurance Program, a semi-annual distribution of three different sample matrices containing various combinations of radionuclides are analyzed. The results of our participation in this program for 1999 are published in ANL-00/2.³¹

Many factors enter into an overall quality assurance program other than the analytical quality control discussed above. Representative sampling is of prime importance. Appropriate sampling protocols are followed for each type of sampling being conducted. Water samples are pre-treated in a manner designed to maintain the integrity of the analytical constituent. For example, samples for trace radionuclide analyses are acidified immediately after collection to prevent hydrolytic loss of metal ions and filtered to reduce leaching from suspended solids.

The monitoring wells are sampled using the protocols listed in the Resource Conservation and Recovery Act (RCRA) Ground Water Monitoring Technical Enforcement Guidance Document. The volume of water in the casing is determined by measuring the water depth from the surface and the depth to the bottom of the well. This latter measurement also determines whether siltation has occurred that might restrict water movement in the screen area. For those wells in the glacial drift that do not recharge rapidly, the well is emptied and the volume removed is compared to the calculated volume. In most cases, these volumes are nearly identical. The well is then sampled by bailing with a Teflon bailer. If samples for

parameters such as priority pollutants are collected, field parameters for these samples (pH, specific conductance, redox potential, and temperature) are measured per well volume while purging. Most samples are collected for radiological analyses only. For samples in the porous, saturated zone which recharge rapidly, three well volumes are purged using submersible pumps. If field parameters are measured, samples are collected as soon as these readings stabilize. All samples are placed in precleaned bottles, labeled, and preserved. All field measurement and sampling equipment is cleaned by field rinsing with Type II deionized water. The samples are transferred to the analytical laboratory along with a computer floppy disk which generates a one-page list of all samples. This list acts as the chain-of-custody transfer document.

6.2 Applicable Standards

The standard that is relevant to this study is the DOE Order 5400.5 which established a dose limit of 100 mrem/y.²¹ The dose limit and dose calculation methodology are applicable to all media: surface water, deep holes, boreholes, and drinking water. The EPA drinking water standard²² is not applicable to the picnic wells since they do not meet the definition of a public water system. However, the EPA standard of 20 nCi/L for tritium may be useful for some comparison purposes.

6.3 Analytical Methods

The analytical methods used to obtain the data in this report are the same as those used in ANL-00/2.³¹

Distribution for ANL-01/1

Internal

R. Bouie
A. B. Cohen
T. M. Davis
A. T. Fracaro
N. W. Golchert (25)
J. E. Helt
M. A. Kamiya
R. G. Kolzow
W. D. Luck
D. A. Milinko
L. P. Moos
J. Sedlet
J. L. Tucker
R. A. Wynveen
TIS File

External

ANL-E Library
ANL-W Library, AW-IS
S. Countiss, DOE Carlsbad Area Office
J. Gilmore, DOE Grand Junction Office (25)
S. L. Heston, DOE Chicago Operations Office (5)
B. J. Quirke, DOE Chicago Operations Office
R. Allen, Illinois Department of Nuclear Safety
J. Barnett, U. S. Environmental Protection Agency, Region V
Bedford Park Public Library
T. Hyde, U. S. Environmental Protection Agency
T. Kelleher, Chicago District, Corps of Engineers
R. LaMort, Cook County Board of Commissioners
Librarian, Illinois Department of Nuclear Safety
A. Martin, Jr., U. S. Geological Survey
Mayor of Willow Springs
M. McMullan, U. S. Environmental Protection Agency, Environmental Review Branch
C. Merenowicz, Forest Preserve District of Cook County
J. R. Nicholas, U. S. Geological Survey
T. Ortciger, Illinois Department of Nuclear Safety
L. Regner, U. S. Environmental Protection Agency, Federal Facility Coordinator
S. Shemanski, Cook County Department of Public Health
C. Smith, Illinois Environmental Protection Agency
D. Weber, Forest Preserve District of Cook County

